Amendments to the Claims

- 1. (Original) A method for treating inflammatory diseases of the external segment or the anterior segment of the eyes, which comprises administering an aqueous eye drop comprising 2-amino-3-(4-bromobenzoyl)phenylacetic acid or its pharmacologically acceptable salt or a hydrate thereof once a day, and maintaining a therapeutically effective concentration of 2-amino-3-(4-bromobenzoyl)phenylacetic acid in the anterior aqueous humor for at least 24 hours after the intraocular administration.
- **2. (Original)** The method according to claim 1, wherein the aqueous eye drop comprising 2-amino-3-(4-bromobenzoyl)phenylacetic acid or its pharmacologically acceptable salt or a hydrate thereof comprises an organic amine or a salt thereof, and the content of the aforementioned organic amine or salt thereof is an amount to allow the octanol-water partition coefficient of 2-amino-3-(4-bromobenzoyl)phenylacetic acid to be 0.7 to 4.
- 3. (Currently amended) The method according to claim 1-or claim 2, wherein the concentration of 2-amino-3-(4-bromobenzoyl)phenylacetic acid or its pharmacologically acceptable salt or a hydrate thereof in the aqueous eye drop is 0.01 to 0.5w/v%.
- **4. (Original)** The method according to claim 2, wherein the organic amine is at least one member selected from the group consisting of an amino acid, an alkanolamine, a diamine, a piperazine, and an aminoalkylsulfonic acid.
- 5. (Currently amended) The method according to claim 2-or claim 4, wherein the organic amine is an amino acid and its concentration is 0.35 to 5w/v%.
- 6. (Currently amended) The method according to claim 2-or claim 4, wherein the organic amine is an alkanolamine and its concentration is 0.15 to 0.95w/v%.

- 7. (Original) The method according to claim 6, wherein the alkanolamine is trometamol.
- **8.** (Currently amended) The method according to claim 2-or claim 4, wherein the organic amine is a diamine and its concentration is 0.05 to 5w/v%.
- 9. (Currently amended) The method according to claim 2-or claim 4, wherein the organic amine is a piperazine and it is contained at a concentration of 0.05 to 5w/v% in the aqueous eye drop.
- 10. (Currently amended) The method according to claim 2-or claim 4, wherein the organic amine is an aminoalkylsulfonic acid and its concentration is 0.05 to 5w/v%.
- 11. (Currently amended) The method according to claim 10, wherein the aminoalkylsulfonic acid is aminoethylsulfonic acid.
- 12. (Original) A method for promoting intraocular penetration of 2-amino-3-(4-bromobenzoyl)phenylacetic acid or its pharmacologically acceptable salt or a hydrate thereof, which comprises administering an aqueous eye drop comprising 2-amino-3-(4-bromobenzoyl)phenylacetic acid or its pharmacologically acceptable salt or a hydrate thereof comprises an organic amine or a salt thereof, wherein the content of said organic amine or salt thereof is an amount to allow the octanol-water partition coefficient of 2-amino-3-(4-bromobenzoyl)phenylacetic acid to be 0.7 to 4.
- 13. (Original) An aqueous eye drop for once a day administration for treating inflammatory diseases of the external segment or the anterior segment of the eyes, characterized in that a therapeutically effective concentration of 2-amino-3-(4-bromobenzoyl)phenylacetic acid or its pharmacologically acceptable salt or a hydrate thereof is maintained in the anterior aqueous humor for at least 24 hours by once a day administration.

- 14. (Original) The aqueous eye drop according to claim 13, wherein the concentration of 2-amino-3-(4-bromobenzoyl)phenylacetic acid or its pharmacologically acceptable salt or a hydrate thereof in the aqueous eye drop is 0.01 to 0.5w/v%.
- 15. (Currently amended) The aqueous eye drop according to claim 13-or elaim-14, wherein the aqueous eye drop comprising 2-amino-3-(4-bromobenzoyl)phenylacetic acid or its pharmacologically acceptable salt or a hydrate thereof comprises an organic amine or a salt thereof, and the content of said organic amine or salt thereof is an amount to allow the octanol-water partition coefficient of 2-amino-3-(4-bromobenzoyl)phenylacetic acid contained in the aqueous eye drop to be 0.7 to 4.
- 16. (Original) The aqueous eye drop according to claim 15, wherein the organic amine is at least one member selected from the group consisting of an amino acid, an alkanolamine, a diamine, a piperazine, and an aminoalkylsulfonic acid.
- 17. (Currently amended) The aqueous eye drop according to claim 15-or elaim 16, wherein the organic amine is an amino acid and its concentration is 0.35 to 5w/v%.
- 18. (Currently amended) The aqueous eye drop according to claim 15-or elaim 16, wherein the organic amine is an alkanolamine and its concentration is 0.15 to 0.95w/v%.
- 19. (Original) The aqueous eye drop according to claim 18, wherein the alkanolamine is trometamol.
- **20.** (Currently amended) The aqueous eye drop according to claim 15-or elaim 16, wherein the organic amine is a diamine and its concentration is 0.05 to 5w/v%.

- **21.** (Currently amended) The aqueous eye drop according to claim 15-or elaim-16, wherein the organic amine is a piperazine and it is contained at a concentration of 0.05 to 5w/v% in the aqueous eye drop.
- **22.** (Currently amended) The aqueous eye drop according to claim 15-or elaim 16, wherein the organic amine is an aminoalkylsulfonic acid and its concentration is 0.05 to 5w/v%.
- **23.** (Original) The aqueous eye drop according to claim 22, wherein the aminoalkylsulfonic acid is aminoethylsulfonic acid.
- 24. (Original) An aqueous eye drop for once a day for treating inflammatory diseases of the external segment or the anterior segment of the eyes, characterized in that a therapeutically effective concentration of 2-amino-3-(4-bromobenzoyl)phenylacetic acid is maintained in the anterior aqueous humor for at least 24 hours by once a day administration of 0.01 to 0.5w/v% 2-amino-3-(4-bromobenzoyl)phenylacetic acid or its pharmacologically acceptable salt or a hydrate thereof and 0.05 to 5w/v% aminoethylsulfonic acid.
- 25. (Currently amended) Use of an organic amine or its salt A method for producing an aqueous eye drop which is to be administered once a day, characterized in that the aqueous eye drop comprising 2-amino-3-(4-bromobenzoyl)phenylacetic acid or its pharmacologically acceptable salt or a hydrate thereof is administered once a day, and a therapeutically effective concentration of 2-amino-3-(4- bromobenzoyl)phenylacetic acid is maintained in the anterior aqueous humor for at least 24 hours by once a day administration to treat inflammatory diseases of the external segment or the anterior segment of the eyes.
- **26.** (Currently amended) The use method according to claim 25, wherein the concentration of 2-amino-3-(4-bromobenzoyl) phenylacetic acid or its pharmacologically acceptable salt or a hydrate thereof in the aqueous eye drop is 0.01 to 0.5w/v%.

- **27.** (Currently amended) The use method according to claim 25-or claim 26, wherein the amount of the organic amine or its salt used is an amount to allow the octanol-water partition coefficient of 2-amino-3-(4-bromobenzoyl)phenylacetic acid to be 0.7 to 4.
- **28.** (Currently amended) The use method according to claim 27, wherein the organic amine is at least one member selected from the group consisting of an amino acid, an alkanolamine, a diamine, a piperazine, and an aminoalkylsulfonic acid.
- 29. (Currently amended) The use method according to claim 27 or claim 28, wherein the organic amine is an amino acid, and its concentration is 0.35 to 5w/v%.
- 30. (Currently amended) The use method according to claim 27-or-claim 28, wherein the organic amine is an alkanolamine, and its concentration is 0.15 to 0.95w/v%.
- 31. (Currently amended) The use method according to claim 30, wherein the alkanolamine is trometamol.
- 32. (Currently amended) The use method according to claim 27-or claim 28, wherein the organic amine is a diamine, and its concentration is 0.05 to 5w/v%.
- 33. (Currently amended) The use method according to claim 27-or-claim 28, wherein the organic amine is a piperazine, and it is contained at a concentration of 0.05 to 5w/v% in the aqueous eye drop.
- 34. (Currently amended) The use method according to claim 27-or claim 28, wherein the organic amine is an aminoalkylsulfonic acid, and its concentration is 0.05 to 5w/v%.

35. (Currently amended) The use method according to claim 34, wherein the aminoalkylsulfonic acid is aminoethylsulfonic acid.